FDA 510(k) Premarket Notification

K812688 QI 1150-1T INFUSION T

CLEARED

Oct 2, 1981
10 days to decision

K812688 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k812688/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 22, 1981
Decision date	Oct 2, 1981
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003